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NEWS 7 Mar 04 SDI PACKAGE for monthly delivery of multifile SDI results
NEWS 8 Mar 24 PATDPAFULL now available on STN
NEWS 9 Mar 24 Additional information for trade-named substances without
structures available in REGISTRY
NEWS 10 Apr 11 Display formats in DGENE enhanced
NEWS 11 Apr 14 MEDLINE Reload
NEWS 12 Apr 17 Polymer searching in REGISTRY enhanced
NEWS 13 Jun 13 Indexing from 1947 to 1956 added to records in CA/CAPLUS
NEWS 14 Apr 21 New current-awareness alert (SDI) frequency in
WPIDS/WPINDEX/WPIX
NEWS 15 Apr 28 RDISCLOSURE now available on STN
NEWS 16 May 05 Pharmacokinetic information and systematic chemical names
added to PHAR
NEWS 17 May 15 MEDLINE file segment of TOXCENTER reloaded
NEWS 18 May 15 Supporter information for ENCOMPAT and ENCOMPLIT updated
NEWS 19 May 19 Simultaneous left and right truncation added to WSCA
NEWS 20 May 19 RAPRA enhanced with new search field, simultaneous left and
right truncation
NEWS 21 Jun 06 Simultaneous left and right truncation added to CBNB
NEWS 22 Jun 06 PASCAL enhanced with additional data
NEWS 23 Jun 20 2003 edition of the FSTA Thesaurus is now available
NEWS 24 Jun 25 HSDB has been reloaded
NEWS 25 Jul 16 Data from 1960-1976 added to RDISCLOSURE
NEWS 26 Jul 21 Identification of STN records implemented
NEWS 27 Jul 21 Polymer class term count added to REGISTRY
NEWS 28 Jul 22 INPADOC: Basic index (/BI) enhanced; Simultaneous Left and
Right Truncation available
NEWS 29 AUG 05 New pricing for EUROPATFULL and PCTFULL effective
August 1, 2003

NEWS EXPRESS April 4 CURRENT WINDOWS VERSION IS V6.01a, CURRENT
MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP),
AND CURRENT DISCOVER FILE IS DATED 01 APRIL 2003
NEWS HOURS STN Operating Hours Plus Help Desk Availability
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STRUCTURE FILE UPDATES: 8 AUG 2003 HIGHEST RN 563538-18-1
DICTIONARY FILE UPDATES: 8 AUG 2003 HIGHEST RN 563538-18-1

TSCA INFORMATION NOW CURRENT THROUGH JANUARY 6, 2003

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Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP
PROPERTIES for more information. See STN Note 27, Searching Properties
in the CAS Registry File, for complete details:
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=> s aeararraaaraara/sqep
 0 AEARRRAAARAARA/SQEP
 97158 SQL=18
L1 0 AEARRRAAARAARA/SQEP
 (AEARRRAAARAARA/SQEP AND SQL=18)

=> s AEARRRAAARAARA/SQsP
L2 0 AEARRRAAARAARA/SQSP

=> s araarraaaraara/sqsp
L3 0 ARAARRAARAARARAE/SQSP

=> s ARAARRAARAARARAE/SQeP
 0 ARAARRAARAARARAE/SQEP
 97158 SQL=18
L4 0 ARAARRAARAARARAE/SQEP
 (ARAARRAARAARARAE/SQEP AND SQL=18)

=> s arraaraarrarae
L5 0 ARRAARAARRARAE

=> s arraaraarrarae/sqsp or arraaraarrarae/sqep

3 ARRAARAAARRARAEA/SQSP
 2 ARRAARAAARRARAEA/SQEP
 40731-SQL=16 - - -
 2 ARRAARAAARRARAEA/SQEP
 (ARRAARAAARRARAEA/SQEP AND SQL=16)
 L6 3 ARRAARAAARRARAEA/SQSP OR ARRAARAAARRARAEA/SQEP

=> s crraaraaarraraea

L7 0 CRRAARAAARRARAEA

=> s CRRAARAAARRARAEA/sqep or CRRAARAAARRARAEA/sqsp

1 CRRAARAAARRARAEA/SQEP
 40731 SQL=16
 1 CRRAARAAARRARAEA/SQEP
 (CRRAARAAARRARAEA/SQEP AND SQL=16)
 1 CRRAARAAARRARAEA/SQSP
 L8 1 CRRAARAAARRARAEA/SQEP OR CRRAARAAARRARAEA/SQSP

=> s l6 or l8

L9 4 L6 OR L8

=> file caplus

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FULL ESTIMATED COST		142.04	142.25

FILE 'CAPLUS' ENTERED AT 12:39:47 ON 11 AUG 2003
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FILE COVERS 1907 - 11 Aug 2003 VOL 139 ISS 7
 FILE LAST UPDATED: 10 Aug 2003 (20030810/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s l9

L10 2 L9

=> d l10 1-2 all

L10 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN
 AN 2001:178433 CAPLUS
 DN 134:217197
 TI Heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects
 IN Harris, Robert B.; Sobel, Michael
 PA Commonwealth Biotechnologies, Inc., USA
 SO U.S., 36 pp., Cont.-in-part of U.S. 5,877,153.

CODEN: USXXAM

DT Patent

LA English

IC ICM A61K038-00

NCL 514013000

CC 1-8 (Pharmacology)

FAN.CNT 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 6200955	B1	20010313	US 1998-166930	19981006
US 5877153	A	19990302	US 1996-660592	19960611
CA 2257614	AA	19971218	CA 1997-2257614	19970603
EP 999219	A2	20000510	EP 1999-119514	19991001
EP 999219	A3	20000524		

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO

JP 2000136202 A2 20000516 JP 1999-285185 19991006

PRAI US 1996-660592 A2 19960611

US 1998-166930 A 19981006

AB Heparin antagonist peptides are disclosed. The heparin-binding peptides of the invention specifically neutralize heparin's conventional anticoagulant properties without causing deleterious hemodynamic side-effects or exacerbation of the proliferative vascular response to injury. More specifically, the heparin-binding compds. of the invention are short-duration drugs to be used in elective or emergency situations which can safely and specifically neutralize heparin's conventional anticoagulant properties without causing deleterious hemodynamic side-effects or exacerbation of the proliferative vascular response to injury.

ST peptide heparin binding anticoagulant neutralization

IT Cell proliferation

Drug delivery systems

Hemostatics

Molecular association

Molecular modeling

Thermodynamics

(heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT Oligosaccharides, biological studies

RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(pentasaccharides, heparin pentasaccharide unit structure; heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT Blood vessel

(smooth muscle; heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT Drug delivery systems

(topical; heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 109319-16-6

RL: BSU (Biological study, unclassified); BIOL (Biological study) (K569-I580 peptide; heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 329358-77-2 329358-78-3 329358-79-4 329358-80-7 329358-81-8 329358-82-9 329358-83-0

RL: PRP (Properties)

(Unclaimed; heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 9000-94-6, Antithrombin III

RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)

(and K121-A134 peptide; heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 9005-49-6, Heparin, biological studies
 RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
 (heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 141509-39-9P 267419-40-9P 267419-42-1P 267419-44-3P 267419-45-4P 268539-71-5P
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 9004-10-8, Insulin, biological studies 268539-65-7 268539-69-1
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 9002-05-5, Blood coagulation factor Xa
 RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
 (heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

IT 329688-19-9
 RL: PRP (Properties)
 (unclaimed protein sequence; heparin binding peptides for neutralization of heparin anticoagulant activity without deleterious side-effects)

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

(1) Anon; WO 9513083 1995 CAPLUS
 (2) Anon; WO 9747312 1997 CAPLUS
 (3) Harris; US 5877153 1999 CAPLUS

L10 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN

AN 1998:13853 CAPLUS

DN 128:93197

TI Novel heparin-binding peptides

IN Harris, Robert B.; Sobel, Michael

PA Commonwealth Biotechnologies, Inc., USA

SO PCT Int. Appl., 61 pp.

CODEN: PIXXD2

DT Patent

LA English

IC ICM A61K038-00

CC 63-6 (Pharmaceuticals)

Section cross-reference(s): 1, 34

FAN.CNT 3

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI	WO 9747312	A1	19971218	WO 1997-US9037 19970603
	W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, UZ, VN, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW: GH, KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG			

US 5877153 A 19990302 US 1996-660592 19960611
CA 2257614 AA 19971218 CA 1997-2257614 19970603
AU-9732167 A1 19980107 AU 1997-32167 19970603
EP 907368 A1 19990414 EP 1997-927795 19970603
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, FI

JP 2001503018 T2 20010306 JP 1998-501628 19970603
PRAI US 1996-660592 A 19960611
WO 1997-US9037 W 19970603

AB The present invention provides heparin antagonist peptides. The heparin-binding peptides of the present invention specifically neutralize heparin's conventional anticoagulant properties without causing deleterious hemodynamic side effects or exacerbation of the proliferative vascular response to injury. More specifically, the heparin-binding compds. of the present invention are short-duration drugs to be used in elective or emergency situations which can safely and specifically neutralize heparin's conventional anticoagulant properties without causing deleterious hemodynamic side effects or exacerbation of the proliferative vascular response to injury.

ST heparin inhibitor peptide sequence

IT Anticoagulants

(inhibitors of; novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

IT Protein sequences

(novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

IT Peptides, biological studies

RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
(novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

IT Drug delivery systems

(topical; novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

IT Amino acids, biological studies

RL: BOC (Biological occurrence); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); OCCU (Occurrence); USES (Uses)
(D-; novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

IT 200888-86-4D, acetyl and succinyl derivs. 200888-87-5D, acetyl and succinyl derivs. 200888-88-6D, acetyl and succinyl derivs.

RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
(novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

IT 9005-49-6, Heparin, biological studies

RL: BPR (Biological process); BSU (Biological study, unclassified); BIOL (Biological study); PROC (Process)
(novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

IT 9004-10-8, Insulin, biological studies

RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)
(novel heparin-binding peptides for neutralizing the anticoagulant effect of heparin)

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Last logoff: 09aug03 11:19:34

Logon file405 11aug03 11:32:10

*** ANNOUNCEMENT ***

--File 654 - US published applications from March 15, 2001 to the present are now online. Please see HELP NEWS 654 for details.

--File 581 - The 2003 annual reload of Population Demographics is complete. Please see Help News581 for details.

--File 990 - NewsRoom now contains February 2003 to current records.
File 992 - NewsRoom 2003 archive has been newly created and contains records from January 2003. The oldest months's records roll out of File 990 and into File 992 on the first weekend of each month.
To search all 2003 records BEGIN 990, 992, or B NEWS2003, a new OneSearch category.

--Connect Time joins DialUnits as pricing options on Dialog.
See HELP CONNECT for information.

--SourceOne patents are now delivered to your email inbox as PDF replacing TIFF delivery. See HELP SOURCE1 for more information.

--Important news for public and academic libraries. See HELP LIBRARY for more information.

--Important Notice to Freelance Authors--
See HELP FREELANCE for more information

NEW FILES RELEASED

***World News Connection (File 985)

***Dialog NewsRoom - 2003 Archive (File 992)

***TRADEMARKSCAN-Czech Republic (File 680)

***TRADEMARKSCAN-Hungary (File 681)

***TRADEMARKSCAN-Poland (File 682)

UPDATING RESUMED

RELOADED

***Population Demographics -(File 581)

***CLAIMS Citation (Files 220-222)

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>>> of new databases, price changes, etc. <<<

* * * * See HELP NEWS 225 for information on new search prefixes
and display codes

SYSTEM:HOME

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*** DIALOG HOMEBASE(SM) Main Menu ***

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2. Database, Rates, & Command Descriptions
3. Help in Choosing Databases for Your Topic
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Enter an option number to view information or to connect to an online
service. Enter a BEGIN command plus a file number to search a database
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? b 410

11aug03 11:32:11 User268147 Session D130.1

\$0.00 0.155 DialUnits FileHomeBase

\$0.00 Estimated cost FileHomeBase

\$0.00 Estimated cost this search

\$0.00 Estimated total session cost 0.155 DialUnits

File 410:Chronolog(R) 1981-2003/Aug

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HIGHLIGHT set on as "

? b 5, 34, 155, 172

11aug03 11:32:23 User268147 Session D130.2

\$0.00 0.071 DialUnits File410

\$0.00 Estimated cost File410

\$0.04 TELNET

\$0.04 Estimated cost this search

\$0.04 Estimated total session cost 0.226 DialUnits

SYSTEM:OS - DIALOG OneSearch

File 5:Biosis Previews(R) 1969-2003/Aug W1

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File 34:SciSearch(R) Cited Ref Sci 1990-2003/Aug W1

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File 155:MEDLINE(R) 1966-2003/Aug W2

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*File 155: Medline has been reloaded and accession numbers have changed. Please see HELP NEWS 155.

File 172:EMBASE Alert 2003/Aug W2

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7/9/1 (Item 1 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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11032797 BIOSIS NO.: 199799653942

High molecular weight kininogen peptides inhibit the formation of kallikrein on endothelial cell surfaces and subsequent urokinase-dependent plasmin formation.

AUTHOR: Lin Yingzhang; Harris Robert B; Yan Wuyi; McCrae Keith R; Zhang Hong; Colman Robert W(a

AUTHOR ADDRESS: (a)Sol Sherry Thrombosis Res. Cent., Temple Univ. Sch. Med., 3400 N. Broad St., Philadelphia, PA 19**USA

JOURNAL: Blood 90 (2):p690-697 1997

ISSN: 0006-4971

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: A sequence of 31 amino acids (S565-K595) in domain 6 of the light chain of high molecular weight kininogen (HK) has previously been shown to be responsible for the binding of plasma prekallikrein (PK) or kallikrein. To find effective peptides that might block binding between HK and PK on cell surfaces, a new series of synthetic peptides has now been prepared that incorporates portions of this binding domain sequence. For mapping the minimal sequence within HK, these new peptides were tested for their ability to compete with HK for binding PK in a cell-free system and on human umbilical vein endothelial cells (HUVEC). In the former, at pH 7.4, the kds for binding between kallikrein and either D567-K595, S565-P594, D567-S593, or D567-T591 were all similar to that for the binding of S565-K595 (0.2 to 0.4 μ -mol/L), but those for the binding of D568-K595, W569-K595, and D567-P589 were an order of magnitude greater (k_d = 2 to 5 μ -mol/L). D567-S586, the shortest chain length of the N- and C-terminal truncation sequences tested, does not effectively compete with kininogen for kallikrein binding (k_d = 100 μ -mol/L). These results imply that D567-T591, a 25-residue peptide (HK25c), contains sufficient structural information for binding kallikrein in solution. D567-T591 also is the minimum structural sequence to block binding of kallikrein to HUVEC-bound HK (IC-50 = 50 nmol/L) and to inhibit PK activation to kallikrein on the cell surface (IC-50 = 80 nmol/L). In addition, D567-T591 also inhibits the generation of kallikrein-activated urokinase, which activates plasminogen to plasmin (IC-50 = 100 nmol/L). Thus, HK-derived peptides may be useful compounds for modulating excessive fibrinolysis and hypotension in sepsis and multiple trauma.

REGISTRY NUMBERS: 9001-01-8: KALLIKREIN; 9039-53-6: UROKINASE; 9001-90-5: PLASMIN

DESCRIPTORS:

MAJOR CONCEPTS: Blood and Lymphatics (Transport and Circulation);

Cardiovascular System (Transport and Circulation); Cell Biology;
Enzymology (Biochemistry and Molecular Biophysics)
BIOSYSTEMATIC NAMES: Hominidae--Primates, Mammalia, Vertebrata, Chordata,
Animalia
ORGANISMS: human (Hominidae)
BIOSYSTEMATIC CLASSIFICATION (SUPER TAXA): animals; chordates; humans;
mammals; primates; vertebrates
CHEMICALS & BIOCHEMICALS: KALLIKREIN; UROKINASE; PLASMIN
MISCELLANEOUS TERMS: Research Article; BIOCHEMISTRY AND BIOPHYSICS;
BLOOD AND LYMPHATICS; CELL SURFACE; CIRCULATORY SYSTEM; FORMATION;
FORMATION INHIBITION; HIGH MOLECULAR WEIGHT; KALLIKREIN; KININOGEN
PEPTIDES; UMBILICAL VEIN ENDOTHELIAL CELL; UROKINASE-DEPENDENT PLASMIN
CONCEPT CODES:
02508 Cytology and Cytochemistry-Human
10808 Enzymes-Physiological Studies
14504 Cardiovascular System-Physiology and Biochemistry
15004 Blood, Blood-Forming Organs and Body Fluids-Blood Cell Studies
10064 Biochemical Studies-Proteins, Peptides and Amino Acids
BIOSYSTEMATIC CODES:
86215 Hominidae

7/9/2 (Item 2 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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11019571 BIOSIS NO.: 199799640716
Physical and biological significance of peptide sequences mediating the
interaction between high molecular weight kininogen and plasma
prekallikrein.
AUTHOR: Colman Robert W(a); Lin Yingzhang; Yan Wuyi; McCrae Keith R; Shenoy
Shilpa S; Harris Robert B
AUTHOR ADDRESS: (a)Sol Sherry Thrombosis Res. Cent., Temple Univ. Sch.
Med., Philadelphia, PA 19140**USA
JOURNAL: Immunopharmacology 36 (2-3):p193-200 1997
ISSN: 0162-3109
RECORD TYPE: Abstract
LANGUAGE: English

ABSTRACT: HK31 (S565-K595) has previously been shown to encompass the
binding domain for plasma prekallikrein (PK) within domain 6 of high
molecular weight kininogen (HK). The complementary binding domain for HK
within PK is mapped to PK56 (F56-G86), in the Apple 1 domain and to PK266
(K266-C295) in the Apple 4 domain. Isothermal titration calorimetry
demonstrated that either PK peptide binds to HK31 in 1:1 stoichiometry.
Binding of the alternate PK peptide into a ternary complex is facilitated
nearly 2-fold. Fluorescence emission spectroscopy revealed that only the
binding of PK56 caused a limited decrease in intrinsic tryptophane
fluorescence emission intensity of HK31. We conclude that the two PK
peptides bind to the HK peptide at different sites. To map the minimal
sequence within HK31, truncated new peptides were tested for their
ability to compete with HK for binding PK in a cell-free system.
D567-T591, a 25-residue peptide which contains sufficient structural
information for binding kallikrein in solution, blocked the binding of
kallikrein to HK bound to endothelial cells and inhibited PK activation
to kallikrein and the generation of kallikrein-activated urokinase on
endothelial cell surfaces. HK-derived peptides could modulate excessive
fibrinolysis and hypotension in sepsis and multiple trauma.

REGISTRY NUMBERS: 9055-02-1: PREKALLIKREIN; 9001-01-8: KALLIKREIN;
9039-53-6: UROKINASE; 9001-90-5: PLASMIN
DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Cardiovascular System (Transport and Circulation); Cell Biology; Endocrine System (Chemical Coordination and Homeostasis); Enzymology (Biochemistry and Molecular Biophysics); Pharmacology

CHEMICALS & BIOCHEMICALS: PREKALLIKREIN; KALLIKREIN; UROKINASE; PLASMIN

MISCELLANEOUS TERMS: Research Article; BINDING DOMAIN; CARDIOVASCULAR SYSTEM; CIRCULATORY SYSTEM; ENDOCRINE SYSTEM; ENDOTHELIAL CELL; ENZYMOLOGY; FIBRINOLYSIS; HIGH-MOLECULAR WEIGHT; HK31; KALLIKREIN; KININOGEN; PEPTIDE SEQUENCES; PHARMACODYNAMICS; PLASMA; PLASMIN; PREKALLIKREIN; S565-K595; UROKINASE

CONCEPT CODES:

02506 Cytology and Cytochemistry-Animal
10064 Biochemical Studies-Proteins, Peptides and Amino Acids
10806 Enzymes-Chemical and Physical
14504 Cardiovascular System-Physiology and Biochemistry
17020 Endocrine System-Neuroendocrinology (1972-)
22010 Pharmacology-Cardiovascular System

? s (sepsis or septic?) and "heparin binding"

91872 SEPSIS

108584 SEPTIC?

154 HEPARIN BINDING

S8 0 (SEPSIS OR SEPTIC?) AND "HEPARIN BINDING"

? s lps and "heparin binding"

78158 LPS

154 HEPARIN BINDING

S9 2 LPS AND "HEPARIN BINDING"

? type s9/full/all

9/9/1 (Item 1 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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13292390 BIOSIS NO.: 200100499539

Biologically active peptides from functional domains of bactericidal/permeability-increasing protein and uses thereof.

AUTHOR: Little Roger G II(a)

AUTHOR ADDRESS: (a)Benicia, CA**USA

JOURNAL: Official Gazette of the United States Patent and Trademark Office

Patents 1246 (2):pNo Pagination May 8, 2001

MEDIUM: e-file

PATENT NUMBER: US 6228834 PATENT DATE GRANTED: May 08, 2001 20010508

PATENT ASSIGNEE: Xoma Corporation PATENT COUNTRY: USA

ISSN: 0098-1133

DOCUMENT TYPE: Patent

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: The present invention provides peptides having an amino acid sequence that is the amino acid sequence of a human bactericidal/permeability-increasing protein (BPI) functional domain or a subsequence thereof, and variants of the sequence or subsequence thereof, having at least one of the BPI biological activities, such as heparin binding, heparin neutralization, LPS binding, LPS neutralization or bactericidal activity. The invention provides peptides and pharmaceutical compositions of such peptides for a variety of therapeutic uses.

DESCRIPTORS:

MAJOR CONCEPTS: Pharmacology

CHEMICALS & BIOCHEMICALS: bactericidal/permeability-increasing protein --human; peptides--biologically active, pharmaceutical

MISCELLANEOUS TERMS: BPI amino acid sequence; BPI biological

activities--LPS binding, LPS neutralization, bactericidal activity, heparin binding, heparin neutralization

9/9/2 (Item 2 from file: 5)

DIALOG(R)File 5:Biosis Previews(R)

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Partial biochemical characterization and purification of IgG2b inducing factor as a new cytokine from synovial fluid of patients with rheumatoid arthritis.

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DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: Rheumatoid arthritis synovial fluid (RA-SF) contains a novel biological activity, which selectively induces IgG2b antibody production in lipopolysaccharide (LPS)-activated mouse spleen cells in vitro and in vivo. Our previous studies have shown that this activity is not functionally identical to other well-known cytokines and interleukins. In this study we demonstrate the partial purification and biochemical characterization of the IgG2b inducing activity in RA-SF. Biochemical characterization revealed that the IgG2b inducing activity in RA-SF has the following properties: it is a protein, sensitive to pH gt 11 and lt 4, which is precipitated by 50% of saturated ammonium sulphate and has a molecular weight of 50-70 kDa; it binds to Cibacron-blue and heparin and its activity is not mediated by immunoglobulins or immune complexes, which are present in RA-SF. Biochemical characteristics of the IgG2b inducing activity also differ from other cytokines and interleukins. The term IgG2b inducing factor is proposed for this novel activity.

REGISTRY NUMBERS: 7783-20-2: AMMONIUM SULFATE

DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Blood and Lymphatics (Transport and Circulation); Clinical Immunology (Human Medicine, Medical Sciences); Endocrine System (Chemical Coordination and Homeostasis); Methods and Techniques; Pathology; Physiology; Skeletal System (Movement and Support)

BIOSYSTEMATIC NAMES: Hominidae--Primates, Mammalia, Vertebrata, Chordata, Animalia

ORGANISMS: Hominidae (Hominidae)

BIOSYSTEMATIC CLASSIFICATION (SUPER TAXA): animals; chordates; humans; mammals; primates; vertebrates

CHEMICALS & BIOCHEMICALS: AMMONIUM SULFATE

MISCELLANEOUS TERMS: AMMONIUM SULFATE PRECIPITATION; ANALYTICAL METHOD; CIBACRON-BLUE BINDING; GEL FILTRATION; HEPARIN BINDING; IMMUNOGLOBULIN G2B; MOLECULAR MASS; PH EFFECT

CONCEPT CODES:

10054 Biochemical Methods-Proteins, Peptides and Amino Acids

10064 Biochemical Studies-Proteins, Peptides and Amino Acids

10506 Biophysics-Molecular Properties and Macromolecules

12508 Pathology, General and Miscellaneous-Inflammation and Inflammatory Disease

15008 Blood, Blood-Forming Organs and Body Fluids-Lymphatic Tissue and Reticuloendothelial System

15010 Blood, Blood-Forming Organs and Body Fluids-Other Body Fluids

17002 Endocrine System-General
 18006 Bones, Joints, Fasciae, Connective and Adipose Tissue-Pathology
 34508 Immunology and Immunochemistry-Immunopathology, Tissue Immunology
 10050 Biochemical Methods-General
 10060 Biochemical Studies-General
 10068 Biochemical Studies-Carbohydrates
 10504 Biophysics-General Biophysical Techniques
 BIOSYSTEMATIC CODES:
 86215 Hominidae
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S2	54	AU='HARRIS ROBERT' OR AU='HARRIS ROBERT B'
S3	36	AU='WOLZ R L' OR AU='WOLZ RL' OR AU='WOLZ RUSSELL' OR AU='WOLZ RUSSELL L'
S4	15	AU='WOLZ G' OR AU='WOLZ G S' OR AU='WOLZ GABRIELLA'
S5	105	S2 OR S3 OR S4
S6	108586	S5 AND SEPSIS OR SEPTIC?
S7	2	S5 AND (SEPSIS OR SEPTIC?)
S8	0	(SEPSIS OR SEPTIC?) AND "HEPARIN BINDING"
S9	2	LPS AND "HEPARIN BINDING"
S10	0	AEARARRAAARAARAARA